Exhibit H

Drugs@FDA: FDA-Approved Drugs

Home (index.cfm) | Previous Page

New Drug Application (NDA): 205029
Company: BELCHER

Products on NDA 205029

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Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	TE Code	RLD
EPINEPHRINE	EPINEPHRINE	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	SOLUTION;INTRAVENOUS, INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS	Prescription	None	Yes

Showing 1 to 1 of 1 entries

Approval Date(s) and History, Letters, Labels, Reviews for NDA 205029

Original Approvals or Tentative Approvals

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Action Date	Submission	Action Type	Submission Classification	Review Priority; Orphan Status	Letters, Reviews, Labels, Pati
07/29/2014	ORIG-1	Approval	Type 7 - Drug Already Marketed without Approved NDA	STANDARD	**Label (PDF) (https://www.accessda Letter (PDF) (https://www.accessda Review (https://www.accessdata.fd**

Showing 1 to 1 of 1 entries

Supplements

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Action Date	Submission	Supplement Categories or Approval Type	Letters, Reviews, Labels, Patient Package Insert
05/18/2016	SUPPL-4	Labeling-Package Insert	**Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/label/2016/205 Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/appletter/2016/**
02/11/2016	SUPPL-2	Efficacy-New Indication	**Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/label/2016/205 Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/appletter/2016/**
12/03/2015	SUPPL-3	Manufacturing (CMC)	
10/23/2015	SUPPL-1	Efficacy-New Indication	**Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/205 Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/appletter/2015/**

Showing 1 to 4 of 4 entries

Labels for NDA 205029